84 Dietz, Holz, Dauer, et al

- 10 Reiber JH, van Eldik Helleman P, Visser Akkerman N, et al. Variabilities in measurement of coronary arterial dimensions resulting from variations in cineframe selection. Cathet Cardiovasc Diagn 1988;14:221–8.
- Storger H. Diffuse in-stent restenosis. J Interv Cardiol 2001;14:587–96.
 Goldberg SL, Loussararian A, De Gregorio J, et al. Predictors of diffuse and
- aggressive intra-stent restenosis. J Am Coll Cardiol 2001;37:1019–25.

 13 Colombo A, De Gregorio J, Moussa I, et al. Intravascular ultrasound-guided percutaneous transluminal coronary angioplasty with provisional spot stenting for treatment of long coronary lesions. J Am Coll Cardiol 2001;38:1427–33.
- 14 Holmes DR Jr, Kip KE, Yeh W, et al. Long-term analysis of conventional coronary balloon angioplasty and an initial "stent-like" result. The NHLBI PTCA Registry. J Am Coll Cardiol 1998;32:590–5.
- 15 Agema WR, Monraats PS, Zwinderman AH, et al. Current PTCA practice and clinical outcomes in the Netherlands: the real world in the pre-drug-eluting stent era. Eur Heart J 2004;25:1163–70.
- 16 Unverdorben M, Sippel B, Degenhardt R, et al. Comparison of a silicon carbide-coated stent versus a noncoated stent in human beings: the Tenax versus Nir stent study's long-term outcome. Am Heart J 2003;145:e17.
- 17 Serruys PW, Jsselmuiden S, Hout B, et al. Direct stenting with the Bx VELOCITY balloon-expandable stent mounted on the Raptor rapid exchange delivery

- system versus predilatation in a European randomized trial: the VELVET trial. Int J Cardiovasc Intervent 2003;5:17–26.
- 18 Jorgensen E, Kelbaek H, Helqvist S, et al. Low restenosis rate of the NIR coronary stent: results of the Danish multicenter stent study (DANSTENT)—a randomized trial comparing a first-generation stent with a second-generation stent. Am Heart J 2003;145:e5.
- 19 Lemos PA, Serruys PW, van Domburg RT, et al. Unrestricted utilization of sirolimus-eluting stents compared with conventional bare stent implantation in the "real world": the rapamycin-eluting stent evaluated at Rotterdam Cardiology Hospital (RESEARCH) registry. Circulation 2004:109:190-5.
- 20 Kornowski R, Mehran R, Hong MK, et al. Procedural results and late clinical outcomes after placement of three or more stents in single coronary lesions. Circulation 1998;97:1355–61.
- 21 Pache J, Kastrati A, Mehilli J, et al. Intracoronary stenting and angiographic results: strut thickness effect on restenosis outcome (ISAR-STEREO-2) trial. J Am Coll Cardiol 2003;41:1283–8.
- 22 Rittersma SZ, de Winter RJ, Koch KT, et al. Impact of strut thickness on late luminal loss after coronary artery stent placement. Am J Cardiol 2004:93:477-80.

IMAGES IN CARDIOLOGY.....

doi: 10.1136/hrt.2005.064642

Percutaneous removal of embolised vegetation from left main coronary artery

55 year old man presented with acute central chest pain two weeks after undergoing redo aortic valve replacement for *Streptococcus mitis* prosthetic endocarditis. His past history included cadaveric renal transplantation for adult polycystic kidney disease. He was receiving intravenous antibiotics at presentation and was afebrile with a normal white count. His ECG is shown in the upper panel.

Primary percutaneous coronary intervention (PCI) was planned. Diagnostic angiography via the right radial demonstrated a filling defect in the left main stem not present at preoperative angiography (panel A)

Emergency bypass surgery was considered to be of prohibitively high risk. In view of the likely haemodynamic instability a percutaneous left ventricular assist device (PVAD) (TandemHeart) was inserted (panel B, arrow). The 21 French venous cannula which was passed across the atrial septum (after predilatation of a patent foramen ovale) and the 17 French femoral artery cannula allowed a flow of approximately 3 l/min. Following this the vegetation was aspirated using an Angiojet device. The filling defect was successfully removed but the left main ostium appeared compromised (panel C). A 3.5×20 mm Taxus stent was deployed in the left main ostium, which was then post-dilated with a 4×20 Quantum non-compliant balloon. An excellent angiographic result (panel D) was obtained with no evidence of major distal embolism. The PVAD was removed after 48 hours. Over the next week, the patient's ejection fraction improved from 25% to 45% and he was discharged home on day 10 after intervention.

> W H T Smith M R Wolff T Kohmoto whs@medicine.wisc.edu



